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APPLICATION FOR LETTERS PATENT

for

**METHOD OF FABRICATING A CONTACT STRUCTURE
HAVING A COMPOSITE BARRIER LAYER BETWEEN
A PLATINUM LAYER AND A POLYSILICON PLUG**

Inventors:
Paul J. Schuele
Pierre C. Fazan

Attorneys:
James R. Duzan
Registration No. 28,393
TRASKBRITT, PC
P.O. Box 2550
Salt Lake City, Utah 84110
(801) 532-1922

TITLE OF THE INVENTION

METHOD OF FABRICATING A CONTACT STRUCTURE HAVING A COMPOSITE BARRIER LAYER BETWEEN A PLATINUM LAYER AND A POLYSILICON PLUG

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application is a continuation of application Serial No. 09/944,436, filed August 30, 2001, pending, which is a continuation of application Serial No. 09/535,050, filed March 24, 2000, now U.S. Patent 6,313,031, issued November 6, 2001, which is a continuation of application Serial No. 08/290,655, filed August 15, 1994, now U.S. Patent 6,093,615, issued July 25, 2000.

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

[0002] This invention was made with Government support under Contract No. MDA972-93-C-0033 awarded by Advanced Research Projects Agency (ARPA). The Government has certain rights in this invention.

BACKGROUND OF THE INVENTION

[0003] Field of the Invention: This invention relates to the manufacture of integrated circuits and, more specifically, to barrier films which are employed to prevent atomic and ionic migration between two adjacent layers which must be conductively coupled.

[0004] State of the Art: For a particular design of an ultra-high-density dynamic random access memory, the integration of a capacitor having a high-permittivity dielectric layer in a CMOS process flow requires that contact be made between a platinum lower capacitor plate and a polycrystalline silicon (polysilicon) plug which makes contact to the storage node junction of the cell access transistor. The dielectric layer, which may be a perovskite oxide such as barium strontium titanate, is deposited at high temperatures in ambient oxygen.

[0005] There are two problems inherent to the dielectric deposition process. The first problem is that the high temperature required for the deposition will initiate a silicidation

reaction between platinum and silicon, thus consuming the platinum capacitor plate and contaminating the capacitor with silicon. It is, therefore, necessary to utilize an electrically conductive diffusion barrier between the platinum and the polysilicon. The second problem is that oxygen will diffuse through the platinum layer and form an insulative silicon dioxide layer between the polysilicon plug and the platinum layer. In such a case, the lower plate of the capacitor will not be in electrical contact with the storage-node junction. Thus, the diffusion barrier must also be impermeable to oxygen.

[0006] Reactively sputtered titanium nitride has been used extensively as a barrier layer in integrated circuits. However, reactively sputtered titanium nitride has a crystalline structure and does not exhibit good step coverage, particularly in deep contact openings. The crystal boundaries associated with such a structure tend to promote ionic and atomic migration. Given this fact, the polysilicon plugs will not be sufficiently protected from reaction with the platinum capacitor plate or with oxygen.

[0007] Titanium nitride deposited via low-pressure chemical vapor deposition (LPCVD) using tetrakis-dimethylamidotitanium or related compounds as the sole precursor is an amorphous material, having no crystal structure and, therefore, no crystal grain boundaries to facilitate atomic and ionic diffusion. However, titanium nitride films deposited via LPCVD have a high carbon content. From X-Ray spectrographic analysis, it appears that some of the carbon atoms have reacted with the titanium to form titanium carbide. The balance of the carbon atoms appears to be unreacted, but trapped, nevertheless, in the titanium nitride/titanium carbide matrix. It is hypothesized that a crystalline structure fails to form because the presence of carbon interferes with crystal nucleation. The presence of carbon, though likely responsible for the amorphous structure of the film (a beneficial quality), is also problematic, as it greatly increases the sheet resistance of the film. In addition, when the carbon-containing films are subjected to high temperatures in the presence of oxygen, the films become perforated and, hence, worthless as barrier films. The perforation phenomenon may be caused by the explosive formation of carbon dioxide gas within the film.

[0008] What is needed is a way to combine the beneficial qualities of both reactively-sputtered titanium nitride with those of titanium nitride deposited via LPCVD.

BRIEF SUMMARY OF THE INVENTION

[0009] This invention is a process for forming an effective titanium nitride barrier layer between the upper surface of a polysilicon plug and a platinum lower capacitor plate in a dynamic random access memory. The memory, which is fabricated on a silicon wafer or substrate, has insulated gate field effect cell access transistors with polysilicon gates which may have a layer of a refractory metal silicide (e.g., tungsten silicide) on their upper surfaces to lower sheet resistance. The transistors are covered with a planarized, thick, flowably deposited dielectric layer such as borophosphosilicate glass (BPSG). A via or contact opening is etched through the thick dielectric layer (the term "thick" is employed merely to distinguish this dielectric layer from the subsequently deposited capacitive dielectric layer) to the storage-node junction of each access transistor, and each of these openings is filled with a polysilicon plug. To begin the barrier layer formation process, the upper surface of each polysilicon plug is recessed at least 1000Å below the upper surface of the thick dielectric layer using a selective polysilicon etch. Using a collimated sputter source, a titanium layer having a thickness of 100-500Å is deposited over the surface of the in-process wafer, thus covering the upper surfaces of the polysilicon plugs. A layer of amorphous titanium carbonitride having a thickness of 100-300Å is then deposited via low-pressure chemical vapor deposition. This is followed by the deposition of a reactively sputtered titanium nitride layer having a thickness of 1000-2000Å. The wafer is then planarized using chemical mechanical polishing (CMP) to remove the titanium, titanium carbonitride and titanium nitride, except that which is in the recesses on top of the silicon plugs. The wafer is then annealed in nitrogen using a rapid thermal anneal system to react the titanium layer with the silicon on the upper surfaces of the plugs to form titanium silicide. The anneal step also repairs damage sustained by the titanium nitride layer during the CMP step. A platinum layer is then deposited and patterned to form lower capacitor electrodes which are electrically coupled to the polysilicon plugs through the titanium silicide, titanium nitride and titanium carbonitride layers. A high permittivity capacitor dielectric layer is then deposited and the capacitors are completed by depositing an upper cell plate layer. The memory is then completed in a conventional manner.

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS

[0010] FIG. 1, a cross-sectional view of a portion of an in-process semiconductor wafer, depicts a portion of a dynamic random access memory (DRAM) array following field oxidation, wordline formation, source/drain implants, deposition of a thick dielectric layer, and polysilicon plug formation;

[0011] FIG. 2 is a cross-sectional view of the portion of the in-process DRAM array of FIG. 1 following an etch selective for polysilicon which has recessed the upper surface of each plug below the upper surface of the thick dielectric layer;

[0012] FIG. 3 is a cross-sectional view of the portion of the in-process DRAM array of FIG. 2 following the deposition of a platinum metal layer;

[0013] FIG. 4 is a cross-sectional view of the portion of the in-process DRAM array of FIG. 3 following the formation of a titanium carbonitride layer via low-pressure chemical vapor deposition and the subsequent deposition of a titanium nitride layer via reactive sputtering;

[0014] FIG. 5 is a cross-sectional view of the portion of the in-process DRAM array of FIG. 4 following a planarization step; and

[0015] FIG. 6 is a cross-sectional view of the portion of the in-process DRAM array of FIG. 5 following the deposition and patterning of a platinum metal layer and the subsequent deposition of a high-permittivity dielectric layer.

DETAILED DESCRIPTION OF THE INVENTION

[0016] Referring now to FIG. 1, a portion of an in-process dynamic random access memory (DRAM) array is shown following a field oxidation step which formed field oxide regions 11, the deposition and patterning of a polysilicon layer to form wordlines WL1-WL4, source/drain implants which have formed storage-node junctions 12 and bit line contact junctions 13, the deposition of a thick, flowable dielectric layer (e.g., borophosphosilicate glass) 14 which covers substrate-superadjacent topography such as wordlines and field oxide regions, the formation of contact openings 15 in the thick dielectric layer 14 which expose the storage-node junctions 12, filling the contact openings 15 with polysilicon material, and planarization of the wafer to form polysilicon plugs 16.

[0017] Referring now to FIG. 2, the polysilicon plugs 16 have been recessed within the contact openings 15 by performing an etch step that is selective for polysilicon, thus forming a well 21 in the thick dielectric layer 14 above each plug. The etch is allowed to proceed until the plugs are recessed at least 1000Å below the upper surface of the thick dielectric layer 14.

[0018] Referring now to FIG. 3, a titanium metal layer 31 is deposited on the upper surface of the wafer to a thickness of 100-500Å using a collimated sputter source.

[0019] Referring now to FIG. 4, a titanium carbonitride layer 41 having a thickness of 100-300Å is deposited on the upper surface of the wafer via low-pressure chemical vapor deposition (LPCVD). Although the compound is referred to as titanium carbonitride (represented by the chemical formula TiC_xN_y), the stoichiometry of the compound is variable, depending on the conditions under which it is deposited. Its primary constituents are titanium and nitrogen, with the ratio of nitrogen to carbon being within a range of 5:1 to 20:1 for the process disclosed herein. X-Ray spectrographic analysis indicates that in these films, some of the carbon atoms have reacted with the titanium to form titanium carbide. The balance of the carbon atoms appear to be unreacted, but trapped, nevertheless, in the titanium nitride/titanium carbide matrix. It is hypothesized that a crystalline structure fails to form because the presence of carbon interferes with crystal nucleation.

[0020] Deposition of the titanium carbonitride layer 41 takes place in a low-pressure chamber (i.e. a chamber in which pressure has been reduced to between 0.1 and 100 Torr prior to deposition), and utilizes a metal-organic tetrakis-dialkylamido-titanium compound as the sole precursor. Any noble gas, as well as nitrogen or hydrogen, or a mixture of two or more of the foregoing, may be used as a carrier for the precursor compound. The wafer is heated to a temperature within a range of 200-600° C. Precursor molecules which contact the heated wafer are pyrolyzed to form titanium nitride containing variable amounts of carbon impurities, which deposits as a highly conformal film on the wafer. Although the carbon impurities present in the deposited films dramatically increase the sheet resistivity of the film, this increase in resistivity is relatively insignificant due to the relative thinness of the deposited layer. The carbon content of the barrier film may be minimized by utilizing tetrakis-dimethylamido-titanium, $Ti(NMe_2)_4$, as

the precursor, rather than compounds, such as tetrakis-diethylamido-titanium or tetrakis-dibutylamido-titanium, which contain a higher percentage of carbon by weight.

[0021] Still referring to FIG. 4, the deposited titanium carbonitride layer 41 demonstrates excellent step coverage, a high degree of conformality, and an acceptable level of resistivity. Subsequent to the deposition of the titanium carbonitride layer 41, a titanium nitride layer 42 having a thickness of 1000-2000Å is deposited on top of the titanium carbonitride layer 41 via reactive sputtering. Titanium nitride layers deposited via reactive sputtering normally do not exhibit good step coverage. Thus, most of the deposition occurs only on horizontal surfaces.

[0022] Referring now to FIG. 5, the wafer is planarized to remove the titanium, titanium carbonitride and titanium nitride, except that which is in the recesses on top of the polysilicon plugs 16.

[0023] Referring now to FIG. 6, the wafer has been subjected to an anneal step using a rapid thermal anneal system to react the titanium metal layer 31 (FIG. 3) with the silicon on the upper surfaces of the polysilicon plugs 16 to form titanium silicide layer 61. If the silicidation reaction is not allowed to proceed to completion, there will be a titanium metal layer on top of the titanium silicide layer 61. The anneal step also repairs damage sustained by the titanium nitride layer 42 during the CMP step. Subsequent to the anneal step, a platinum metal layer is deposited on the upper surface of the wafer and patterned to form lower capacitor electrodes 62. A high permittivity capacitor dielectric layer 63 is then deposited on the upper surface of the wafer. The capacitor dielectric layer 63, which may be a perovskite oxide such as barium strontium titanate or strontium titanate, is deposited at high temperatures in ambient oxygen. The titanium nitride layer 42 of the bi-layer barrier film that has been fabricated prevents oxygen from diffusing to the titanium carbonitride layer 41, where it would attack free carbon atoms and cause perforations in the barrier layer. Likewise, the titanium carbonitride layer 41, because it is an amorphous material without crystal grain boundaries, is able to prevent the migration of silicon atoms to the titanium nitride layer 42 through which it would be able to diffuse to reach the overlying platinum lower capacitor electrodes 62. The cell capacitors are completed by depositing an upper cell plate layer (not shown). The memory is then completed in a conventional manner.

[0024] Although only a single embodiment of the invention has been described in detail, it will be obvious to those having ordinary skill in the art of integrated circuit manufacture that changes and modifications may be made thereto without departing from the spirit and the scope of the invention as hereinafter claimed. For example, the invention may also be practiced in the context of a dynamic random access memory array having a buried bit line architecture, which bit lines (also known as digit lines) are formed after word line formation, but before cell capacitor formation. Thus, the general process flow depicted is meant to be only illustrative and not limiting.